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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁷ : C07D 239/52, 239/56, 239/46, 239/36, A61K 31/505		A1	(11) International Publication Number: WO 00/03998
			(43) International Publication Date: 27 January 2000 (27.01.00)
<p>(21) International Application Number: PCT/EP99/05134</p> <p>(22) International Filing Date: 19 July 1999 (19.07.99)</p> <p>(30) Priority Data: CA98A000015 17 July 1998 (17.07.98) IT</p> <p>(71) Applicant (for all designated States except US): NOVIRIO PHARMACEUTICALS LIMITED [-/-]; Walker Secretaries, Walker House, Grand Cayman (KY).</p> <p>(72) Inventors; and</p> <p>(75) Inventors/Applicants (for US only): LA COLLA, Paolo [IT/IT]; 5a, Strada, 11, Poggio dei Pini, I-09012 Capoterra (IT). ARTICO, Marino [IT/IT]; Via Edgardo Negri, 64, I-00128 Roma (IT).</p> <p>(74) Agents: MODIANO, Guido et al.; Modiano, Josif, Pisanty & Staub, Baaderstrasse 3, D-80469 München (DE).</p>		<p>(81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).</p> <p>Published With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</p>	
<p>(54) Title: SUBSTITUTED 6-BENZYL-4-OXOPYRIMIDINES, PROCESS FOR THEIR PREPARATION AND PHARMACEUTICAL COMPOSITIONS CONTAINING THEM</p>			
<p>(57) Abstract</p> <p>The invention concerns novel substituted 6-benzyl-4-oxypyrimidines of general formula (A). These compounds inhibit reverse transcriptase encoded by human immunodeficiency virus (HIV) or pharmaceutically acceptable salts thereof, and find their application in the prevention and treatment of HIV infection and the treatment of the resulting acquired immune deficiency syndrome (AIDS). Pharmaceutical compositions containing the compounds and a method of use of the present compounds and other agents for the treatment of AIDS and viral infection by HIV are also envisaged.</p>			
		<p style="text-align: right;">(A)</p>	